### AMENDMENTS TO THE CLAIMS

This listing of claims will replace all prior versions, and listings, of claims in the application:

- 1-7. (canceled)
- 8 (withdrawn, currently amended) The <u>An</u> amyloid-targeting imaging agent of claim 1, of the formula.

$$A_t - A_{lnk} - A_{lab}$$
 (I)

wherein At is of formula

$$Q^{b} = \begin{bmatrix} - & * \\ - & X \end{bmatrix}_{2}$$
 (IV)

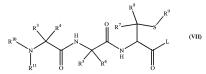
wherein Y is an anionic group at physiological pH;  $Q^b$  is a carrier molecule;  $X^+$  is a cationic group; and  $n^2$  is 2;  $A_{bb}$  is a linker moiety; and  $A_{bb}$  is a labeling moiety<del>an integer selected such that the biodistribution of</del>

the targeting moiety for the intended target site is not prevented while maintaining activity of the targeting moiety.

- (withdrawn) The amyloid-targeting imaging agent of claim 8, wherein Y is a sulfonate group.
- (withdrawn) The amyloid-targeting imaging agent of claim 8, wherein Y is a sulfate or thiosulfate group.
- (withdrawn) The amyloid-targeting imaging agent of claim 8, wherein Y is a tetrazole group.
- 12-20. (canceled)
- (currently amended) The amyloid-targeting imaging agent of <u>claim 31 or claim 32-claim</u>
   wherein A<sub>bab</sub> includes a radionuclide selected from <sup>90ar</sup>Tc, <sup>99</sup>Tc, <sup>64</sup>Cu, <sup>67</sup>Cu, <sup>97</sup>Ru, <sup>109</sup>Pd.

<sup>186</sup>Re, <sup>188</sup>Re, <sup>111</sup>In, <sup>113</sup>mIn, <sup>153</sup>Gd, <sup>90</sup>Y, <sup>153</sup>Sm, <sup>166</sup>Ho, <sup>198</sup>Au, <sup>199</sup>Au, <sup>90</sup>Sr, <sup>89</sup>Sr, <sup>105</sup>Rh, <sup>201</sup>Tl, <sup>51</sup>Cr, <sup>67</sup>Ga, <sup>57</sup>Co, <sup>60</sup>Co, <sup>123</sup>L, <sup>125</sup>L, <sup>131</sup>Lor, <sup>18</sup>F.

- (currently amended) The amyloid-targeting imaging agent of <u>claim 31 or claim 32 claims</u>
   +, wherein A<sub>lab</sub> includes a radionuclide selected from the group consisting of Tc and Re.
- (currently amended) The amyloid-targeting imaging agent of claim 31 or claim 32 claim
   wherein A<sub>bb</sub> is a metal chelate of a radioactive or paramagnetic metal ion.
- (withdrawn, currently amended) The amyloid-targeting imaging agent of claim 31 or claim 32 claim 1, wherein Abh comprises a chelating ligand of the formula



where  $R^{10}$  is a linear or branched, saturated or unsaturated  $C_{14}$  alkylene group interrupted by one or two heteroatoms;  $R^{11}$  is H or  $R^{10}$ , or  $R^{10}$  and  $R^{11}$  taken together, form a 5-to 8-membered saturated or unsaturated heterocyclic ring optionally substituted with one or more of halogen, hydroxyl, amino, carboxyl, oxo,  $C_{1\cdot4}$  alkyl, aryl, or C(O)R groups;  $R^3$ ,  $R^4$ ,  $R^5$  and  $R^6$  are independently H, carboxyl,  $C_{1\cdot4}$  alkyl, an alpha carbon side chain of a D- or L-amino acid other than proline, or C(O)R;  $R^7$  and  $R^8$  are independently H, carboxyl, amino,  $C_{1\cdot4}$  alkyl,  $C_{1\cdot4}$  alkyl;  $R^9$  is H or a sulfur protecting group; and L is hydroxyl, alkoxy, an amino acid residue, or a linking group.

# 25-30. (canceled)

31. (currently amended) The An amyloid-targeting imaging agent of claim 1, of the formula

$$A_1 - A_{lnk} - A_{lab}$$
 (I)

where z is 0 or 1; At is an amyloid targeting moiety of the formula

where

 $R^1$  is an <u>unsubstituted or substituted  $C_{1-20}$  alkyl,  $C_{2-20}$  alkenyl, hydroxyalkyl, or a single-ring aromatic group;</u>

 $R^2$  is a-an unsubstituted or substituted  $C_{1.20}$  alkyl,  $C_{2.20}$  alkenyl, hydroxyalkyl, a single-ring aromatic group, or a hydrogen atom, or  $R^1$  and  $R^2$ , taken together with the nitrogen to which they are attached, form a heterocyclic group-which is a fused ring structure;

T is an alkylene group of the formula –(CH<sub>2</sub>)<sub>n</sub>-wherein n is an integer from 2-12; Y is SO<sub>2</sub>X, and X is a cationic group;

Ank is a linker moiety; and Anh is a labeling moiety.

32. (currently amended) The An amyloid-targeting imaging agent of elaim 1, of the formula

$$A_t - A_{lnk} \rightarrow A_{lab}$$
 (I)

where z is 0 or 1; At is an amyloid targeting moiety of the formula

where

 $R^{1}$  is a  $C_{5}$ - $C_{18}$  <u>unsubstituted or substituted</u> alkyl, <del>hydroxyalkyl</del> or <u>an unsubstituted</u> or <u>substituted</u> single-ring aromatic group;

 $R^2$  is a hydrogen atom or [[an]] a  $C_{1-8}$  alkyl group;

T is an alkylene group of the formula  $-(CH_2)_{h}$ - wherein n is an integer from 2-12; Y is  $SO_3X$ , and X is a cationic group;

Alik is a linker moiety; and Alab is a labeling moiety.

33. (original) The An amyloid-targeting imaging agent of claim 1, of the formula

$$A_t - A_{lnk} - A_{lab}$$
 (I)

where A, is of the formula

$$\begin{bmatrix} R^1 & & & \\ & & & \\ & & & \\ R^2 & & & \\ \end{bmatrix}_{D}^{(T)} (T) (Y)_{s}$$

#### where

- R<sup>1</sup> is an alkyl, an alkenyl, or an aromatic group;
- R<sup>2</sup> is a hydrogen atom, an alkyl group, or an aromatic group, or R<sup>1</sup> and R<sup>2</sup>, taken together, form a heterocyclic group which is a fused ring structure;
- Z and Q are each independently a carbonyl (C=O), thiocarbonyl (C=S), sulfonyl (SO<sub>2</sub>), or sulfoxide (S=O) group;
- k is 1 and m is 0 or 1;
- p and s are each 1;
- T is an alkylene group of the formula -(CH<sub>2</sub>)<sub>n</sub>- wherein n is an integer from 2-12;
- Y is SO<sub>3</sub>X, and X is a cationic group.
- (canceled)
- (withdrawn) The amyloid-targeting imaging agent of claim 33, wherein said A<sub>t</sub> is
  selected from the group consisting of 3-acetylamino-1-propanesulfonic acid, 3benzoylamino-1-propanesulfonic acid, and 2-acrylamido-2-methyl-1-propanesulfonic
  acid.
- (withdrawn) The amyloid-targeting imaging agent of claim 33, wherein said At is selected from the group consisting of 3-phthalimido-1-propanesulfonic acid, N-(3sulfopropy)saccharin and 4-phthalimido-1-butanesulfonic acid.
- 37. (withdrawn) The amyloid-targeting imaging agent of claim 32, wherein said A<sub>t</sub> is selected from the group consisting of 3-phenylamino-1-propanesulfonic acid, 3-(4-pyridylamino)]-1-propanesulfonic acid, 3-(benzylamino)-1-propanesulfonic acid, 2 deoxy-2-(3-sulfopropyl)amino-D-glucose, 3-[(-3,4-dimethyl-1-adamantyl)-amino]-1-propanesulfonic acid, 3-[(-3,5-dimethyl-1-adamantyl)-amino]-1-propanesulfonic acid, 3-(2-hydroxyethyl)amino-1-propanesulfonic acid, 3-(3-hydroxy-1-propyl)amino-1-propanesulfonic acid, 3-

[(d,l)-1-hydroxy-2-propyllamino-1-propanesulfonic acid, 3-(4-hydroxy-1-butyl)amino-1propanesulfonic acid, 3-(5-hydrox-1-pentyl)amino-1-propanesulfonic acid, 3-(6-hydroxy-1-hexyl)amino-1-propanesulfonic acid. 3-(4-hydroxyphenyl)amino-1-propanesulfonic acid. (+)-3-[(S)-2-hydroxy-1-propyllamino-1-propanesulfonic acid. (+)-3-[(S)-1hydroxy-2-propyllamino-1-propanesulfonic acid. (-)-3-f(R)-1-hydroxy-2-propyllamino-1-propanesulfonic acid. (+)-3-f(S)-1-hydroxy-2butyllamino-1-propanesulfonic acid. (-)-3-[(R)-1-hydroxy-2-butyllamino-1propanesulfonic acid, 3-[(dl)-5-hydroxy-2-pentyl]amino-1-propanesulfonic acid, 3-[(dl)-6-hvdroxy-2-hexyllamino-1-propanesulfonic acid, 3-(1-hvdroxymethyl-1cyclopentyl)amino-1-propanesulfonic acid, 3-amylamino-1-propanesulfonic acid, 3hexylamino-1-propanesulfonic acid, 3-heptylamino-1-propanesulfonic acid, 3octylamino-1-propanesulfonic acid, 3-nonylamino-1-propanesulfonic acid, 3-decylamino-1-propanesulfonic acid, 3-undecylamino-1-propanesulfonic acid, 3-dodecylamino-1propanesulfonic acid, 3-tridecylamino-1-propanesulfonic acid, 3-tetradecylamino-1propanesulfonic acid, 3-hexadecylamino-1-propanesulfonic acid, and 3-octadecylamino-1-propanesulfonic acid.

### 38-42. (canceled)

- (currently amended) A kit for preparing a radiopharmaceutical preparation, said kit comprising:
  - an amyloid-targeting imaging agent of claim 31 or claim 32 -claim-1;
  - a reducing agent;
  - a buffering agent;
  - a transchelating agent, and

instructions for the preparation and use of the radiopharmaceutical in the imaging of amyloid or an amyloid-related condition.

# 44-50. (canceled)

 (withdrawn, currently amended) A method of diagnostic medical imaging of an amyloidassociated disease comprising the steps of administering to a patient a pharmaceutical composition according to claim [[1]] 31 or claim 32 and then imaging said patient.

 (withdrawn) The method of diagnostic medical imaging according to claim 51 wherein A<sub>bb</sub> of said pharmaceutical composition is a radiopharmaceutical.

- (withdrawn) The method of diagnostic medical imaging according to claim 51 wherein A<sub>kb</sub> of said pharmaceutical composition is a metal chelate.
- (withdrawn) The method of diagnostic medical imaging according to claim 53 wherein said metal chelate is gadolinium-DTPA, gadolinium-DOTA, or gadolinium-DO3A.
- (withdrawn) The method of diagnostic medical imaging according to claim 53 wherein said metal chelate is a chelate of <sup>99m</sup>Tc or <sup>111</sup>In.
- (withdrawn) The method of diagnostic medical imaging according to claim 51 wherein said imaging step is ultrasound imaging.
- (withdrawn) The method of claim 105, wherein said imaging step is radionuclide imaging.
- 58. (withdrawn) The method of claim 57, wherein said imaging step is SPECT imaging.
- (withdrawn) The method of claim 105, wherein said imaging step is magnetic resonance imaging.
- 60. (withdrawn) The method of claim 105, wherein said imaging step is ultrasound imaging.
- 61. (withdrawn) The method of claim 105, wherein said imaging step is X-ray imaging.
- (withdrawn) The method of claim 105, wherein said imaging step is fluorescence imaging.
- 63-91. (canceled)
- 92. (withdrawn, currently amended) The method of claim [[87]] 105, wherein said A<sub>1</sub> is selected from the group consisting of 3-phenylamino-1-propanesulfonic acid, 3-(4-pyridylamino)]-1-propanesulfonic acid, 3-(benzylamino)-1-propanesulfonic acid, 2-deoxy-2-(3-sulfopropyl)amino-D-glucose, 3-[(-3,4-dimethyl-1-adamantyl)-amino]-1-propanesulfonic acid, 3-(-3,5-dimethyl-1-adamantyl)-amino]-1-propanesulfonic acid, 3-

(2-hydroxyethyl)amino-1-propanesulfonic acid, 3-(3-hydroxy-1-propyl)amino-1propanesulfonic acid, (-)-3-[(R)-2-hydroxy-1-propyl]amino-1-propanesulfonic acid, 3-[(d.l)-1-hydroxy-2-propyllamino-1-propanesulfonic acid. 3-(4-hydroxy-1-butyl)amino-1propanesulfonic acid. 3-(5-hydrox-1-pentyl)amino-1-propanesulfonic acid. 3-(6-hydroxy-1-hexyl)amino-1-propanesulfonic acid, 3-(4-hydroxyphenyl)amino-1-propanesulfonic acid. (+)-3-[(S)-2-hydroxy-1-propyllamino-1-propanesulfonic acid. (+)-3-[(S)-1hydroxy-2-propyllamino-1-propanesulfonic acid. (-)-3-[(R)-1-hydroxy-2-propyllamino-1-propanesulfonic acid. (+)-3-[(S)-1-hydroxy-2-butyllamino-1-propanesulfonic acid. (-)-3-[(R)-1-hydroxy-2-butyllamino-1-propanesulfonic acid, 3-[(d1)-5-hydroxy-2pentyllamino-1-propanesulfonic acid, 3-[(d1)-6-hydroxy-2-hexyl]amino-1propanesulfonic acid, 3-(1-hydroxymethyl-1-cyclopentyl)amino-1-propanesulfonic acid, 3-amylamino-1-propanesulfonic acid, 3-hexylamino-1-propanesulfonic acid, 3heptylamino-1-propanesulfonic acid, 3-octylamino-1-propanesulfonic acid, 3nonylamino-1-propanesulfonic acid, 3-decylamino-1-propanesulfonic acid, 3undecylamino-1-propanesulfonic acid, 3-dodecylamino-1-propanesulfonic acid, 3tridecylamino-1-propanesulfonic acid, 3-tetradecylamino-1-propanesulfonic acid, 3hexadecylamino-1-propanesulfonic acid, and 3-octadecylamino-1-propanesulfonic acid.

### 93-94. (canceled)

95. (withdrawn, currently amended) A method for diagnostic medical imaging of an amyloid-associated disease in a patient, comprising administering to a patient a pharmaceutical composition comprising an amyloid-targeting imaging agent of <u>claim 31 or claim 32</u>, the formula

$$A_{l}$$
  $A_{lnk}$   $A_{lab}$   $A_{lab}$ 

as defined in claim 63 and then imaging the amyloid-targeting imaging agent in said patient..

 (withdrawn) The method of claim 95, wherein A<sub>lab</sub> of said pharmaceutical composition is a radiopharmaceutical.

 (withdrawn) The method of claim 95, wherein A<sub>lab</sub> of said pharmaceutical composition is a metal chelate.

- (withdrawn) The method of claim 95, wherein A<sub>lab</sub> of said pharmaceutical composition is a metal chelate and said imaging step is magnetic resonance imaging or radionuclide imaging.
- (withdrawn) The method of claim 97, wherein said metal chelate is gadolinium-DTPA, gadolinium-DOTA, or gadolinium-DO3A.
- 100. (withdrawn) The method of claim 97, wherein said metal chelate is a chelate of <sup>99m</sup>Tc or <sup>111</sup>In.
- (withdrawn, currently amended) The method of claim [[63]] <u>95</u>, wherein said imaging step is ultrasound imaging.
- 102. (canceled)
- 103. (withdrawn, currently amended) A method for diagnosing an amyloid-related condition in a patient, comprising administering an amyloid-targeting imaging agent according to claim [[1]] 31 or claim 32 to a patient, and imaging said amyloid-targeting imaging agent in said patient to determine the presence of amyloid in said patient, such that the presence or absence of an amyloid-related condition in said patient is determined.
- 104. (withdrawn) The method of claim 103, wherein said amyloid-related condition is selected from the group consisting of Creutzfeld-Jakob Disease (CJD), Kuru, transmissible cerebral amyloidoses (also known as transmissible virus dementias), familial CJD, scrapic, transmissible mink encephalopathy, bovine spongiform encephalopathy (BSE), inflammation-associated amyloid, type II diabetes, primary amyloidosis, feline spongiform encephalopathy, non-transmissible cerebral amyloidosis (e.g., Alzheimer's disease), prion-mediated diseases, dialysis-related amyloidosis, tight chain-related amyloidosis, cerebral amyloid angiopathy, and Alzheimer's disease.
- 105. (withdrawn, currently amended) A method for imaging amyloid deposition in a patient, comprising administering an amyloid-targeting imaging agent according to claim [[1]] 31

- or claim 32 to a patient, and imaging said amyloid-targeting imaging agent in said patient to determine the presence of amyloid in said patient.
- 106 (new) The amyloid-targeting imaging agent of claim 31, wherein n is an integer from 2 to 4.
- 107. (new) The amyloid-targeting imaging agent of claim 31, wherein the heterocyclic group is a six-membered heterocyclic group.
- 108 (new) The amyloid-targeting imaging agent of claim 107, wherein the six-membered heterocyclic group is piperidyl.
- 109. (new) The amyloid-targeting imaging agent of claim 108, wherein A<sub>t</sub> is 4(1-piperidyl)-1-butanesulfonic acid, or a pharmaceutically acceptable salt thereof.
- 110. (new) The amyloid-targeting imaging agent of claim 107, wherein the heterocyclyl is 1,2,3,6-tetrahydropyridyl.
- 111. (new) The amyloid-targeting imaging agent of claim 110, wherein A<sub>t</sub> is 3[1-(1,2,3,6-tetrahydropyridyl)]-1-propanesulfonic acid, or a pharmaceutically acceptable salt thereof.
- 112 (new) The amyloid-targeting imaging agent of the formula in claim 32, wherein n is an integer from 2 to 4.
- 113. (new) The amyloid-targeting imaging agent of claim 32, wherein R<sup>1</sup> is an alkyl group which is substituted with an amino group and a carboxyl group.
- 114 (new) The amyloid-targeting imaging agent of claim 113, wherein R<sup>1</sup> is 5-amino-1-carboxy-1-pentyl; R<sup>2</sup> is hydrogen.
- 115 (new) The amyloid-targeting imaging agent of claim 114, wherein A<sub>t</sub> is a-N-(3-sulfopropyl)-L-lysine, or a pharmaceutically acceptable salt thereof.
- 116 (new) The amyloid-targeting imaging agent of claim 32, wherein R<sup>1</sup> is an alkyl group which is substituted with a hydroxyl group.

117. (new) The amyloid-targeting imaging agent of claim 116, wherein R<sup>1</sup> is 6-hydroxy-1-hexyl.

- 118 (new) The amyloid-targeting imaging agent of claim 117, wherein R<sup>2</sup> is hydrogen.
- (new) The amyloid-targeting imaging agent of claim 117, wherein A<sub>t</sub> is 3-(6-hydroxy-1-hexyl)amino-1-propanesulfonic acid, or a pharmaceutically acceptable salt thereof.
- 120. (new) The amyloid-targeting imaging agent of claim 32, wherein R<sup>1</sup> is an unsubstituted or substituted C<sub>4-7</sub> cycloalkyl group.
- (new) The amyloid-targeting imaging agent of claim 120, wherein the cycloalkyl is substituted with a hydroxymethyl.
- 122 (new) The amyloid-targeting imaging agent of claim 121, wherein the cycloalkyl is cyclopentyl.
- 123. (new) The amyloid-targeting imaging agent of claim 120, wherein R<sup>1</sup> is 1-hydroxy-1-cyclopentyl; R<sup>2</sup> is hydrogen.
- 124 (new) The amyloid-targeting imaging agent of claim 123, wherein A<sub>t</sub> is 3(1-hydroxy-1-cyclopentyl)amino-1-propanesulfonic acid, or a pharmaceutically acceptable salt thereof.
- 125. (New) The amyloid-targeting imaging agent of claim 8, wherein Y is a sulfonate or a sulfate group.
- 126. (New) The amyloid-targeting imaging agent of claim 125, wherein Q<sup>b</sup> is NHCH<sub>2</sub>CH<sub>2</sub>CH<sub>2</sub>-.
- (New) The amyloid-targeting imaging agent of claim 126, wherein At is 3-hydroxy-1propylsulfamic acid O-sulfate or a pharmaceutically acceptable salt thereof.
- 128. (New) The amyloid-targeting imaging agent of claim 9, wherein Q<sup>b</sup> is an alkylene or a piperizine-1.4-bis(alkylene).
- 129. (New) The amyloid-targeting imaging agent of claim 128, wherein A<sub>t</sub> is 1,6-hexanedisulfonic acid or a pharmaceutically acceptable salt.

130. (New) The amyloid-targeting imaging agent of claim 128, wherein A<sub>t</sub> is 1,4 piperizinebis(propanesulfonic acid) or a pharmaceutically acceptable salt.

131. (New) The amyloid-targeting imaging agent of claim 9, where A<sub>t</sub> is Thiazole yellow G.